WHAT IS CLAIMED IS:

1. A compound according to formula I

1300×

Ind-Q-N Z-R1

Ι

wherein

Ind is unsubstituted indol-3-yl, indol-3-yl monosubstituted by OH, OA, CN, Hal, COR^2 or CH_2R^2 , or indol-3-yl polysubstituted by OH, OA, CN, Hal, COR^2 , CH_2R^2 or combinations thereof;

R¹ is benzofuran-5-yl, 2,3-dihydrobenzofuran-5-yl, ehroman-6-yl, chroman-4-on-6-yl, 3-chromen-6-yl or chromen-4-on-6-yl, which in each case is unsubstituted or monosubstituted by CN, CH₂OH, CH₂OA or COR²;

Q is C_mH_{2m} ;

Z is Nor CR3;

A is alkyl having 1-6 C atoms;

Hal is F, Cl, Br or I;

 R^2 is OH, OA, NH_2 , NHA or NA_2 ;

R³ is H, OH or OA; and

m is 2, 3 or 4; or

a physiologically acceptable salt thereof.

- 2. A compound according to claim 1, wherein said compound is:
- (a) 1-[4-(5-methoxyindol-3-yl)butyl]-4-(2-hydroxymethylbenzofuran-5-yl)piperazine or a physiologically acceptable salt thereof;



- (b) 1-[4-(5=carbamoylindol-3-yl)butyl]-4-hydroxy-A-(2-3-dihydrobenzofuran-5-yl)piperidine or a physiologically
 -acceptable salt-thereof;
- (c) 1-[4 (5 carbamoylindol-3-yl)butyl] 4 (2,3-dihydrobenzofuran 5-yl)piperidine or a physiologically-acceptable salt thereof;
- -(d) 1 [4 (5 methoxyindol=3 yl)butyl 4 (2,3-dihydrobenzofuran-5-yl)ptperazine or a physiologically acceptable salt thereof;
 - ethoxycarbonylbenzofuran-5-yl)piperazine or a physiologically acceptable salt thereof;
- (1) 1-[4-(5-cyanoindol-3-yl)butyl]-4-(2-carbamoylbenzofuran-5-yl)piperazine or a physiologically acceptable salt thereof
- (g) 1 [4-(5-methoxyindol-3-yl)butyl]-4-(chroman-6-yl)piperazine or a physiologically acceptable salt thereofy
- (h) 1-[4-(5-hydroxyindol-3-yl)butyl]-4-(chroman-6-yl)piperazine-or-a physiologically acceptable salt-thereof.
 - 3. A compound according to claim 1, wherein Ind is unsubstituted indol-3-yl, indol-3-yl monosubstituted by OH, OA, CN, Hal, COR^2 or CH_2R^2 , or indol-3-yl disubstituted by OH, OA, CN, Hal, COR^2 or CH_2R^2 .
 - 4. A compound according to claim 1, wherein Ind is indol-3-yl monosubstituted in the 5-position by OH, OA, CN, Hal, COR^2 or CH_2R^2 .
 - 5. A compound according to claim 1, wherein Ind is indol-3-yl monosubstituted in the 4-, 6- or 7-position by OH, OA, CN, Hal, COR^2 or CH_2R^2 .
 - 6. A compound according to claim 1, wherein A is methyl or ethyl.

- 7. A compound according to claim 1, wherein R^1 is benzofuran-5-yl, $\frac{2}{3}$ -dihydrobenzofuran-5-yl, chroman-6-yl or chroman-4-on-6-yl which, in each case is unsubstituted or monosubstituted by $-CH_2OH$, $-CONH_2$, $-CO_2A$ or $-CO_2NHA$.
- 8. A compound according to claim 1, wherein Q is $-(CH_2)_4-$.
- 9. A compound according to claim 1, wherein Z is -N-, -C(OH) or -CH-.
- 10. A compound according to claim 1, wherein Ind is indol-3-yl substituted in the 5-position by OH or OA.
- A compound according to claim 1, wherein Ind is indol-3-yl substituted in the 5-position by CONH₂ or CN.
- 12. A compound according to claim 1, wherein Z is N and R¹ is unsubstituted benzofuran-5-yl or benzo-5-yl substituted by CN, CH₂OH, CH₂OA or COR².
- 13. A compound according to claim 1, wherein Z is -CH(OH)-.
- 14. A compound according to claim 1, wherein Z is N and \mathbb{R}^1 is 2,3-dihydrobenzofuran-571.
- 15. A compound according to claim 1, wherein Z is N and \mathbb{R}^1 is chroman-6-yl.
- 16. A compound according to claim 1, wherein Z-is-N-and-R¹ is chromen-4-on-6-yl.
- A pharmaceutical composition comprising a compound according to claim 1 and a pharmaceutically acceptable carrier.

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- 18. A composition according to claim 27, wherein said compound is present in an amount of 0.2-500 mg.
- 19. A method of treating tension, depression, psychosis or side effects associated with the treatment of hypertension, comprising administering a compound according to claim 1.
- 20. A method of treating acromegaly, hypogonadism, secondary amenorrhea, premenstrual syndrome, undesired puerperal lactation, or cerebral disorders, comprising administering a compound according to claim 1.
- 21. A method of treating migraines, comprising administering a compound according to claim 1.
- 22. A method according to claim 21, wherein said compound is administered in a daily dosage of 0.001-0.005 mg/kg of body weight.

